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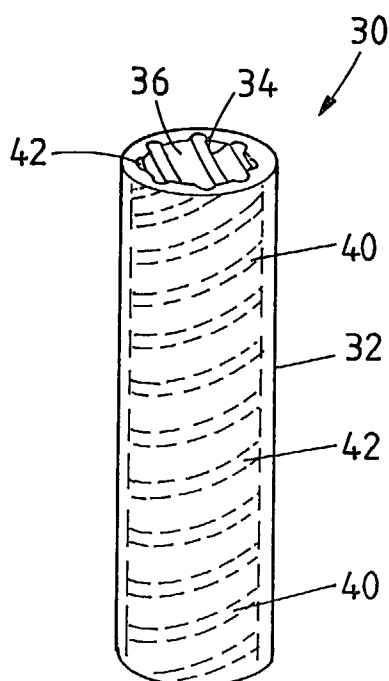
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(54) Title: FLOW ADAPTIVE ASPIRATION TUBING AND DEVICES



(57) Abstract: Aspiration tubing for use in a phacoemulsification system is provided having a modified lumen that enhances resistance of fluid flow through the lumen to reduce post-occlusion surge, the modification comprising recesses or protuberances formed in the interior surface of the lumen, a freely-moving object disposed in the flow path of the lumen, or one or more bends.



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FLOW ADAPTIVE ASPIRATION TUBING AND DEVICES

Field of the Invention

5 **[0001]** The present invention relates generally to apparatus for performing phacoemulsification, in particular to flow adaptive aspiration tubing for use in conjunction with phacoemulsification systems.

10 Background of the Invention

[0002] The crystalline lens of the human eye transmits and focuses light and is located behind the iris attached to the wall of the eye by suspensory ligaments known as the zonules. The lens consists of a
15 more rigid central nucleus surrounded by peripheral cortical material, which has a softer consistency. A fine membrane known as the capsule contains the entire lens.

[0003] Cataract formation refers to a loss of transparency of the crystalline lens of the eye and is a
20 common occurrence with age. This results in a progressive reduction in vision, which can be restored with surgery. Cataract surgery involves removal of the cataractous lens and insertion of a plastic intraocular lens to replace the crystalline lens. Removal of the
25 cataractous lens is accomplished using ultrasonic energy

to fragment and aspirate the lens by a technique known as phacoemulsification.

[0004] During such surgery, a central opening is formed in the anterior portion of the capsule to permit
5 access to the lenticular material. An ultrasonic handpiece, typically including a needle having an outer wall and central lumen, is then inserted, contacted against and caused to fragment the lens. An elastomeric sleeve surrounding the needle provides a conduit
10 irrigating the eye to replace material aspirated through the needle. Once the nuclear material of the lens has been removed with the assistance of ultrasonic energy, softer cortical material may be aspirated with an irrigation/aspiration cannula.

15 [0005] In both phases of the procedure it is important that the anterior chamber is maintained at a positive pressure and constant volume to prevent collapse, so as to prevent trauma to sensitive ocular tissues. Contact with the endothelial cells lining the
20 posterior surface of the cornea or the iris can result in irreparable damage. Even more common is inadvertent contact or aspiration of the posterior capsule, which prevents the escape of the fluid contained in the posterior chamber of the eye known as the vitreous
25 humour. Such inadvertent contact may result in rupture of the posterior capsule membrane.

[0006] Rupture of the posterior capsule and loss of the vitreous humour increases the risk of retinal detachment and cystoid macular oedema after cataract
30 surgery, with subsequent loss of vision. Furthermore if the posterior capsule is disrupted during surgery it may not be feasible to properly place an intraocular lens in in the capsular bag remnant of the original lens, again

resulting in a less favorable outcome than might be anticipated in uncomplicated surgery.

[0007] Maintaining a stable pressure and volume in the anterior chamber when performing phacoemulsification is of paramount concern. Optimal fluid dynamics implies sustaining a stable pressure and volume in the anterior chamber when performing phacoemulsification. Aspiration of fluid from the anterior chamber must be balanced by adequate infusion. The desired state of fluid balance may be summarized in the equation: $F_i = F_o$ - Inflow (F_i) should equal Outflow (F_o). To avoid chamber collapse the pressure in anterior chamber (P_{ac}) also must be greater than atmospheric pressure (P_a) and greater than vitreous pressure (P_v) - $P_{ac} > P_a > P_v$.

[0008] The pressure in the anterior chamber depends on the infusion pressure, which is the difference between the irrigation pressure head (P_i), related to the irrigation bottle height, and the drop in pressure due to resistance to the inflow of irrigation fluid (P_d) - $P_a = P_i - P_d$. The anterior chamber pressure preferably should be maintained at a constant level to avoid alterations in chamber volume, which manifest as an unstable chamber during surgery.

[0009] A conventional apparatus used in cataract surgery includes a console containing a pump system used to generate vacuum and flow as well as the electrical circuitry that provides energy and control for the phacoemulsification handpiece. The pump systems are connected to the phacoemulsification handpiece and irrigation and aspiration cannula by tubing so that fluid and lens material can be aspirated from the eye.

[0010] Several types of pump systems are known for providing aspiration of fluid and lens material during

phacoemulsification and cortical aspiration. The first type are positive fluid displacement pumps, such as a peristaltic pump. In such systems, fluid flow is generated by drawing suction through the tubing and
5 significant vacuum may be achieved if the tubing becomes occluded. In other pump systems, such as a venturi pump, suction is generated in a cassette and the subsequent flow and aspiration of fluid from the eye is related to that preset suction level.

10 [0011] For either pump system, the sequence of removal of nuclear and cortical material is similar. Fluid is aspirated from the anterior chamber via suction applied through the phacoemulsification needle or irrigation/aspiration cannula and the associated
15 aspiration tubing. This suction attracts nuclear or cortical material to the needle or cannula and may result in larger fragments occluding the tip or aspiration port.

[0012] The suction level within the tubing then rises until the negative pressure generated overcomes the
20 resistance of the lenticular material, which is then aspirated down the tubing. This in turn causes a rapid equalization of pressure between the anterior chamber and the rest of the system, with a concomitant rapid increase in flow and drop in chamber pressure. This phenomenon is
25 typically referred to as "post occlusion surge" and may cause a forward movement of the posterior capsule as the chamber pressure and volume fluctuates.

[0013] Vacuum applied by the phacoemulsification handpiece may be modulated by foot pedal control, thereby
30 causing the pump system to respond by venting or equalizing the pressure in the system either to fluid or to air. The venting, however, occurs, some distance from the handpiece and anterior chamber and there is typically

a lag before the vacuum in the tubing is restored to a positive pressure and the pressure in the anterior chamber is restored to the normal resting or unoccluded level.

5 **[0014]** Accordingly, it would be desirable to reduce the surge in flow rate that occurs with rapid fluctuations in vacuum pressure associated with occlusion of the phacoemulsification needle. Such control advantageously could reduce fluctuations in chamber
10 pressure and shorten the time to attain equilibrium pressure, thereby enhancing safety of the surgical procedure and reducing the risk of inadvertent rupture of the posterior capsule.

[0015] One potential method for reducing fluid surges
15 in the aspiration tubing is to reduce the maximum vacuum levels that are generated by the pump system. High vacuum levels, however, are advantageous in capturing fragments of nuclear material so that the fragments may be fractured into smaller pieces. It is therefore
20 desirable to maintain high vacuum levels while reducing the high flow rates associated with surges that occur at those high levels of vacuum.

[0016] Another approach is to increase the resistance in the aspiration tubing by reducing the lumen or
25 increasing the length of the aspiration tubing. While reducing the lumen size may be effective, the internal diameter of typical phacoemulsification tubing is generally about 1.5 mm, and any further reduction in diameter is likely to result in obstruction of the
30 aspiration tubing by lens material. Increasing the tubing length may be accomplished by coiling the tubing to add further hydrodynamic resistance. In both cases, however, the increased resistance of the tubing exists

for all vacuum levels. This is undesirable, as it would be preferable to maintain undiminished aspirational flow rates at low vacuum levels to facilitate attraction of lens fragments prior to occlusion.

5 [0017] In view of the foregoing, it would be desirable to provide a phacoemulsification system including flow adaptive aspiration tubing that automatically increases flow resistance in response to higher flow rates.

10 [0018] It further would be desirable to provide a phacoemulsification system including flow adaptive aspiration tubing that induces turbulent flow at lower flow velocities.

[0019] It would be yet further desirable to provide a
15 phacoemulsification system including flow adaptive aspiration tubing that provides improved anterior chamber stability at higher vacuum levels.

Summary of the Invention

20 [0020] In view of the foregoing, it is an object of the present invention to provide a phacoemulsification system including flow adaptive aspiration tubing that automatically increases flow resistance in response to higher flow rates.

25 [0021] It is another object of the present invention to provide a phacoemulsification system including flow adaptive aspiration tubing that induces turbulent flow at lower flow velocities.

[0022] It is a further object of this invention to
30 provide a phacoemulsification system including flow adaptive aspiration tubing having a modified lumen that defines longitudinal flutes arranged in a spiral pattern.

[0023] It is another object of the present invention to provide a phacoemulsification system including flow adaptive aspiration tubing that provides improved anterior chamber stability at higher vacuum levels.

5 [0024] These and other objects of the present invention are accomplished by providing flow adaptive tubing for use with phacoemulsification systems, wherein the tubing has a lumen surface that enhances turbulent flow through the tubing at relatively low flow
10 velocities.

[0025] In one preferred embodiment the aspiration tubing constructed in accordance with the principles of the present invention comprises topographical features, such as spiral flutes, inwardly projecting protuberances,
15 ridges or recesses, formed on an inner surface of the aspiration tubing. The features may be provided along the entire length of the tubing or only along discrete portions of the length, and may extend around the entire circumference of the lumen or only parts thereof.

20 [0026] In an alternative embodiment the aspiration tubing comprises a freely-moving object, such as a vane or a propeller, disposed within the lumen of the tubing. In a yet further alternative embodiment, the aspiration tubing comprises one or more bends along the length of
25 the tubing that are configured to enhance hydrodynamic resistance at higher flow velocities.

Brief Description of the Drawings

[0027] The above and other objects and advantages of
30 the present invention will be apparent upon consideration of the following detailed description, taken in conjunction with the accompanying drawings, in which like

referenced characters refer to like parts throughout, and in which:

[0028] FIG. 1 is a side-sectional view of a phacoemulsification system suitable for use with the flow adaptive aspiration tubing of the present invention;

[0029] FIG. 2 is a perspective view of a length of aspiration tubing in accordance with a preferred embodiment of the present invention;

[0030] FIG. 3 is a perspective view of a length of aspiration tubing in accordance with another embodiment of the present invention;

[0031] FIG. 4 is a perspective view of a length of aspiration tubing in accordance with an additional embodiment of the present invention;

[0032] FIG. 5 is a perspective view of a length of aspiration tubing in accordance with a further embodiment of the present invention;

[0033] FIG. 6 is a side-sectional view of a length of aspiration tubing in accordance with another embodiment of the present invention including a rotating vane disposed within the aspiration tubing;

[0034] FIG. 7 is a side-sectional view of a length of aspiration tubing in accordance with an additional embodiment of the present invention including a rotating propeller disposed within the aspiration tubing;

[0035] FIG. 8 is a side-sectional view of a length of aspiration tubing including a pair of bends in accordance with a further embodiment of the present invention;

[0036] FIG. 9 is a chart showing the velocity of fluid flow within the aspiration tubing plotted against the drop in pressure due to frictional forces within the aspiration tubing; and

[0037] FIGS. 10A and 10B are perspective and plan views, respectively, of an extrusion die used to extrude the aspiration tubing shown in FIG. 2.

5 Detailed Description of the Invention

Overview of a Preferred Phacoemulsification System

[0038] Referring to FIG. 1, a phacoemulsification system suitable for use with the flow adaptive tubing of the present invention is described. Phacoemulsification
10 system 10 comprises phacoemulsification needle 11 coupled to ultrasonic handpiece 20 and surrounded by elastomeric sleeve 12. Handpiece 20 is typically coupled to a controller (not shown), that causes needle 11 to vibrate at ultrasonic frequencies. When needle is contacted
15 against a cataractous lens, vibration of needle 11 causes the needle to fragment the nuclear material of the lens.

[0039] Irrigation line 13 is coupled in fluid communication between a source of irrigation fluid 14 and handpiece 20 so that irrigation fluid is delivered into
20 the eye via annulus 15 formed between needle 11 and interior surface 16 of elastomeric sleeve 12. Aspiration tubing 17 is coupled in fluid communication between lumen 18 of needle 11 and vacuum source 19, to permit the aspiration of fragmented nuclear material from a
25 patient's eye.

[0040] In operation, ultrasound energy is applied to the nuclear material of the patient's lens by phacoemulsification needle 11. During the phacoemulsification process, irrigation fluid is supplied
30 to the eye from irrigation reservoir 14 via annulus 15 between needle 11 and elastomeric sleeve 12. Simultaneously, fragmented nuclear material is withdrawn through needle lumen 18 and aspiration tubing 17. As

described hereinabove, larger fragments that are caused to engage needle 11 by virtue of suction drawn through lumen 18 may cause surges in flow rates of material aspirated through lumen 18 and wide fluctuations of the pressure within the eye. The aspiration tubing of the present invention is expected to moderate such post-occlusion surges and reduce the resulting pressure fluctuations.

10 ***Description of the Flow Adaptive***

Aspiration Tubing of the Present Invention

[0041] In accordance with the principles of the present invention, phacoemulsification system 10 of FIG. 1 is provided with aspiration tubing having a lumen modified to enhance hydrodynamic resistance to fluid flow at higher flow rates. The modification may take one of a number of forms, and comprise topographic features along the interior surface of the tubing, freely-movable obstructions within the flow path, or one or more flow-redirecting bends.

[0042] Referring now to FIG. 2, a first illustrative embodiment of adaptive aspiration tubing 30 of the present invention is described. Tubing 30 has outer surface 32 and inner surface 34 that defines lumen 36. In accordance with the principles of the present invention, inner surface 34 comprises topographical features 40 that enhance flow resistance at higher flow rates by enhancing turbulent flow within the lumen. Features 40 may be provided along the entire length of the tubing or only along one or more discrete segments of the tubing. In addition, the features may extend entirely around the circumference of tubing 30 as shown in FIG. 2, or may extend for only limited arcs of the circumference.

[0043] In one preferred embodiment, features 40 comprise spiral flutes or recesses 42 formed as grooves in the inner surface of the tubing. Recesses 42 illustratively are disposed at regular intervals along the length of the tubing and around the circumferential periphery of the inner surface. Recesses 42 may have a semi-annular cross-section (as shown in FIG. 2) or alternatively may include a cross-sectional profile of a semi-ellipse, rectangular, triangular, square, diamond or other suitable shape.

[0044] Still referring to FIG. 2, recesses 42 produce turbulent flow within lumen 36 without increasing the likelihood that a large fragment will obstruct the lumen. In particular, recesses 40 enhanced turbulence by increasing the hydrodynamic resistance of fluid passing through the tubing. Advantageously, because flow resistance increases at higher flow rates, risks associated with fluid surge and accompanying pressure fluctuations that could cause collapse or partial collapse of the anterior chamber of the eye are reduced. In addition, features 40 alter the flow pattern within the lumen such that the flow becomes turbulent at lower flow velocities.

[0045] Referring now to FIG. 3, alternative embodiments are described, wherein the topographical features 40 comprise raised protuberances 44, such as ridges or lugs, that project from the inner surface of tubing into lumen 36. Protuberances 44 may be arranged as a series of semi-cylindrical ridges disposed in a spiral pattern at regular intervals along the length of the tubing and around the circumferential periphery of lumen 36. Alternatively, protuberances 44 may extend only partially around the circumference of lumen 36, or

only along selected portions of the tubing length. Like recesses 42, protuberances 44 enhance hydrodynamic resistance within tubing 30 and induce turbulent flow at lower flow velocities and vacuum levels than otherwise encountered in previously known aspiration tubing.

[0046] In FIGS. 4 and 5, further alternative embodiments are depicted, in which topographical features 40 comprise longitudinal recesses 46 (FIG. 4) or longitudinal protuberances 48 (FIG. 5) formed in the inner surface of tubing 30. Longitudinal recesses 46 or protuberances 48 are shown disposed substantially parallel to each other and spaced at regular intervals around the circumferential periphery of lumen 36. Alternatively, recesses 46 or protuberances 48 may extend only partially around the circumference of lumen 36, or only along selected portions of the tubing length.

[0047] As further would be understood by those of skill in the art, any combination of two or more of the above described topographical features may be provided on the inner surface of the aspiration tubing without departing from the scope of the present invention.

[0048] Referring now to FIGS. 6 and 7, and in accordance with another aspect of the present invention, aspiration tubing 30 comprises a freely-moving object 50, illustratively, a vane or propeller, disposed within lumen 36. Object 50 is configured to fit within tubing 30 so that it disrupts laminar flow of fluid within lumen 36 and induces turbulence. In FIGS. 6 and 7, laminar flow is indicated by straight arrow L and turbulent flow is indicated by curved arrow T. The induced turbulence increases hydrodynamic resistance within lumen 36, and is a function of the flow velocity within the tubing.

[0049] By way of example, freely-moving object 50 may comprise vane 56 (FIG. 6) mounted within lumen 36 on hub 58 and having a plurality of arms 60. Alternatively, as depicted in FIG. 7, freely-moving object 50 may comprise
5 propeller 62 having a plurality of projections 63. Propeller 62 is mounted on axle 64 supported on arms 65 within lumen 66 of separate short longitudinal segment 67. Segment 67 includes adapters 68 at either end so that freely-moving object 50 may be used in conjunction
10 with previously-known aspiration tubing. Of course, as will be appreciated by those of skill in the art, freely-moving object 50 may take on other shapes without departing from the scope of the present invention. In addition, flow through the aspiration tubing may be
15 modified using more than one freely-moving object to disrupt laminar flow.

[0050] Referring to FIG. 8, another alternative embodiment of aspiration tubing of the present invention is described, in which aspiration tubing 30 comprises one
20 or more angular bends 70. Bends 70 cause abrupt changes in the direction of fluid flowing within lumen 72, so that an initially laminar flow (as indicated by straight arrow L) becomes turbulent flow (as indicated by curved arrow T) after the fluid passes through the one or more
25 angular bends. Preferably, the bend angles are less than or equal to 90°. As for the embodiment of FIG. 7, the embodiment of FIG. 8 may be implemented as a separate segment of rigid tubing that includes one or more bends, and adapters for coupling the segment in-line with
30 previously-known aspiration tubing.

[0051] As noted above, previously-known aspiration tubing used in cataract surgery has a smooth inner surface. The rate of fluid flow within such tubing is

typically laminar and may be described by the Hagen-Poiseuille equation:

$$Q = P \times \pi \times D^4 / (8 \times l \times v)$$

where Q is the volume flow rate,

5 P is the pressure differential,

D is the cross sectional diameter of the tubing,

l is the length of the restricting diameter, and

v is the viscosity of the fluid.

[0052] The point at which the flow of fluid within the
10 tubing becomes turbulent is determined by the Reynolds number:

$$Re = d \times V \times D / v$$

where Re is Reynolds number,

V is the fluid velocity,

15 D is the cross sectional diameter of the tubing and
v is the fluid viscosity.

Experiments show that flow is likely to be laminar if Reynolds number is less than 2000 (the lower critical Reynolds number) and turbulent if it exceeds 4000 (the
20 upper critical Reynolds number). The nature of flow in tubing is uncertain when the Reynolds number is between 2000 and 4000.

[0053] Increasing the roughness of the inner wall surface of the tubing increases resistance to flow and
25 tends to induce turbulent flow. When fluid flows through a length of modified tubing that lacks a circular cross section (e.g., FIGS. 2-5), the Reynolds number may be calculated using the equivalent diameter d_e where:

$$d_e = 4 \times A / P,$$

30 where A is the cross section of flow area and

P is the cross-sectional wetted perimeter. Altering the profile of tubing therefore promotes turbulent flow.

[0054] When fluid flow is laminar most of the shearing action and friction between layers of fluid takes place away from the wall. The wall surface has relatively little effect on the friction factor (f) which varies

5 inversely with the Reynolds number according to $f = 64 / Re$, where Re is the Reynolds number. By contrast, when flow is turbulent the nature of the inner surface of the tubing has a significant effect on the friction factor, because much of the shearing action and resulting

10 friction takes place near the topographical features. The Relative roughness may be defined as:

$$\epsilon_R = \epsilon / d$$

where ϵ is the height of the surface roughness and d is the tube diameter.

15 The Moody diagram and foregoing formula may be used to calculate the friction factor (f) for fluid flow in tubing: $f = 0.0055 * (1 + (20000 * \epsilon_R + 10^6 / Re)^{1/3})$.

[0055] Accordingly, modifying the inner surface of the tubing to produce an uneven surface causes flow within

20 the tubing to become turbulent at lower velocities and vacuum levels and thus increases the resistance to flow. Turbulent flow can be described by the Bernoulli equation:

$$Q = \text{SQRT}(P * \pi^2 * D^4) / (8 * d)$$

25 where Q is the volume flow rate,
P is the pressure differential,
D is the cross sectional diameter of the tubing,
l is the length of the restricting diameter,
v is the viscosity of the fluid and

30 d is the density of the fluid.

For the same pressure differential that is determined by the vacuum level generated by the vacuum, the flow rate will be less due to the increased resistance produced by

turbulent flow for the same diameter and length of tubing.

[0056] Furthermore, the resistance afforded to fluid flow by the modified aspiration tubing will be relatively unchanged at low vacuum levels but increases proportionally as the vacuum levels rise due to the transition from laminar to turbulent flow (which occurs at lower vacuum level than in previously-known tubing with a smooth internal surface). The drop in pressure due to flow resistance varies as the square of the velocity. Therefore, as depicted in FIG. 9, the drop in pressure plotted against the velocity of fluid flow forms a parabola, wherein a relatively small change in fluid velocity results from a relatively large percentage drop in pressure compared to tubing with a lower frictional resistance and laminar flow.

[0057] Advantageously, the provision of topographical features on the inner surface of the tubing does not reduce the efficiency of the phacoemulsification needle in attracting fragments of nuclear material. When occlusion of the needle tip occurs, the initial rise in vacuum within the tubing is similar to previously-known tubing, but the rise time slows as the vacuum level is increased. When the occlusion resistance is overcome, the immediate flow of fluid due to the negative pressure in the aspiration line will be less than with previously-known tubing. Advantageously, this reduces both the magnitude and duration of post-occlusion surge.

[0058] The present invention provides modified aspiration tubing with increased flow resistance at higher vacuum levels. This provides a more stable anterior chamber pressure and volume than that encountered with previously-known aspiration tubing. In

addition, the aspiration tubing of the present invention enables higher vacuum levels to be employed during phacoemulsification. Higher vacuum levels increase the attraction of nuclear fragments to the needle tip and
5 facilitate the fracture of the fragments into smaller pieces by a second instrument. Moreover, because the cross sectional area and the length of the modified aspiration tubing can be formed to be the same as with previously-known aspiration tubing, the likelihood of
10 obstruction by fragments of lens material is not increased.

[0059] Previously-known aspiration tubing may be formed from extruded plastic materials, wherein the unpolymerized plastic flows out through a conical
15 extrusion nozzle provided with a central die. The shape of the die determines the cross-sectional profile of the internal lumen of the aspiration tubing and is usually cylindrical in form. The speed of extrusion and relationship between the central die and surrounding
20 nozzle determines the wall thickness.

[0060] Referring to FIGS. 10A and 10B, a die suitable for producing aspiration tubing with a fluted spiral inner lumen (as shown in FIG. 2) is described. Die 90 resembles a gear and has central cylindrical portion 92
25 and a plurality of semi-circular protuberances 94 spaced at substantially equal intervals around its outer periphery. By rotating the extruded plastic material as it exits the extrusion nozzle, a spiral pattern of grooves or flutes is formed in the internal surface of
30 the lumen. The speed of rotation determines the pitch of the spiral pattern. Suitable materials for manufacturing the aspiration tubing of the present invention include polyvinyl chloride and silicones. As would be

understood by those of skill in the art, the modified aspiration tubing can be fabricated by other methods than the above-described extruded molding method without departing from the scope of the present invention.

5 [0061] Central cylindrical portion 92 and protuberances 94 preferably are dimensioned so that the cross sectional surface area of the lumen of the fluted spiral tubing is equivalent to that of previously-known tubing having a circular cross section. Further, the arc
10 length of each semi-circle protuberance along the circumference of central cylindrical portion 92 may be calculated as follows:

$$C = 2 \times \pi \times R / 12$$

where R = radius of central cylindrical core.

15 The central angle α subtended by each semi-circle in radians is given by:

$$\alpha = C / R$$

where C = chord length d of each semi-circle:

$$d = 2 \times R \times \sin(\alpha / 2)$$

20 The sagittal height h of the flute is given by:

$$h = R - \sqrt{R^2 - r^2}$$

where r = radius of flute semi-circle

The area S of the segment of produced by intersection of semi-circular protuberance of the flute with the
25 circumference of central cylindrical portion 92 is:

$$S = r^2 \times \text{ACOS}((r - h) / r) - \sqrt{2 \times r \times h - h^2} \times (r - h)$$

The area f of the flute is given by:

$$f = (r^2 \times \pi) / 2 - S$$

The total area T of fluted tube is given by:

30
$$T = R^2 \times \pi - 8 \times f$$

[0062] According to one preferred embodiment, the modified spiral fluted aspiration tubing of the present

invention has an internal diameter of about 1.5 mm and an external diameter of about 3.12 mm. It should be noted that tubing with a lumen having an internal diameter of 1.34 mm and 8 equally-spaced flutes with a radius of
5 0.173 mm will have the same total cross sectional area as previously-known tubing with a lumen having an internal diameter of 1.5 mm. The modified aspiration tubing of the present invention may be constructed with similar connectors attached to the proximal and distal ends as
10 previously-known tubing, and thus have the same overall length.

[0063] Although the invention is described with reference to ultrasound as an energy source to remove the cataract, the modified aspiration tubing also may be used
15 with any method of cataract removal where other energy sources such as laser, sonic, rotary tips, impellers, hydro jet and mechanical methods are used to fragment the cataract and the lens material is removed by aspiration. In addition, it is envisioned that the modified
20 aspiration tubing may be advantageously employed in other medical applications wherein fluids or tissues are aspirated.

[0064] Further, as noted hereinabove, the entire length of aspiration tubing may include the modified
25 lumen. Alternatively, as further noted above, the modified aspiration tubing may be formed as either a flexible or rigid segment that may be inserted into a previously-known aspiration line as a separate device. Further, a three-way tap may be provided within the
30 aspiration line to permit bypass or inclusion of the modified aspiration tubing in the aspiration circuit, and thereby alter the rate of fluid flow within an internal lumen of the aspiration line.

[0065] Although preferred illustrative embodiments of the present invention are described above, it will be evident to one skilled in the art that various changes and modifications may be made without departing from the invention. It is intended in the appended claims to
5 cover all such changes and modifications that fall within the true spirit and scope of the invention.

What Is Claimed Is:

1. Aspiration tubing for use with a phacoemulsification system, the aspiration tubing comprising:

a length of tubing having first and second ends and an interior surface that forms a lumen extending between the first and second ends,

wherein the interior surface further defines a plurality of recesses or protuberances communicating with the lumen that enhance resistance to fluid flow through the lumen.

2. The aspiration tubing of claim 1, wherein the recesses or protuberances are disposed along an entire length of the interior surface between the first and second ends.

3. The aspiration tubing of claim 1 wherein the recesses or protuberances are disposed along one or more segments of the length of the tubing.

4. The aspiration tubing of claim 1, wherein the recesses form spiral flutes.

5. The aspiration tubing of claim 1, wherein the protuberances form annular ridges arranged in a spiral configuration.

6. The aspiration tubing of claim 1, wherein the recesses or protuberances are spaced at regular intervals around a circumference of the lumen.

7. The aspiration tubing of claim 1 further comprising adapters disposed at the first and second ends to couple the aspiration tubing in-line with a phacoemulsification system.

8. Aspiration tubing for use with a phacoemulsification system, the aspiration tubing comprising:

a length of tubing having first and second ends and an interior surface that forms a lumen extending between the first and second ends,

a freely-moving object disposed within the lumen to disrupt the formation of laminar flow near the interior surface and enhance resistance to fluid flow through the lumen.

9. The aspiration tubing of claim 8, wherein the freely-moving object is a vane or a propeller.

10. The aspiration tubing of claim 8 further comprising adapters disposed at the first and second ends to couple the aspiration tubing in-line with a phacoemulsification system.

11. Aspiration tubing for use with a phacoemulsification system, the aspiration tubing comprising:

a length of tubing having first and second ends and an interior surface that forms a lumen extending between the first and second ends,

wherein a portion of the tubing comprises one or more bends having an angle of 90 degrees or less that enhance resistance to fluid flow through the lumen.

12. Apparatus for use with a phacoemulsification system comprising:

a length of tubing having first and second ends and an interior surface that forms a lumen extending between the first and second ends; and

means disposed within the lumen and communicating with the lumen for enhancing resistance to fluid flow through the lumen.

13. The apparatus of claim 12, wherein the means for enhancing resistance to fluid flow through the lumen comprises features formed on the inner surface of the aspiration tubing.

14. The apparatus of claim 13, wherein the features are recesses formed in the interior surface of the aspiration tubing.

15. The apparatus of claim 14, wherein the recesses form spiral flutes.

16. The apparatus of claim 13, wherein the features are protuberances projecting inwardly into the lumen.

17. The apparatus of claim 16, wherein the protuberances form annular ridges arranged in a spiral configuration.

18. The apparatus of claim 13, wherein the features are spaced at regular intervals around a circumference of the lumen.

19. The apparatus of claim 12 further comprising adapters disposed at the first and second ends to couple the aspiration tubing in-line with a phacoemulsification system.

20. The apparatus of claim 12 wherein the means for enhancing resistance to fluid flow through the lumen comprises a freely-moving object disposed within the lumen to disrupt the formation of laminar flow near the interior surface.

21. The apparatus of claim 20, wherein the freely-moving object is a vane or a propeller.

22. The apparatus of claim 12 wherein the means for enhancing resistance to fluid flow through the lumen comprises a portion of the tubing having one or more bends at an angle of 90 degrees or less.

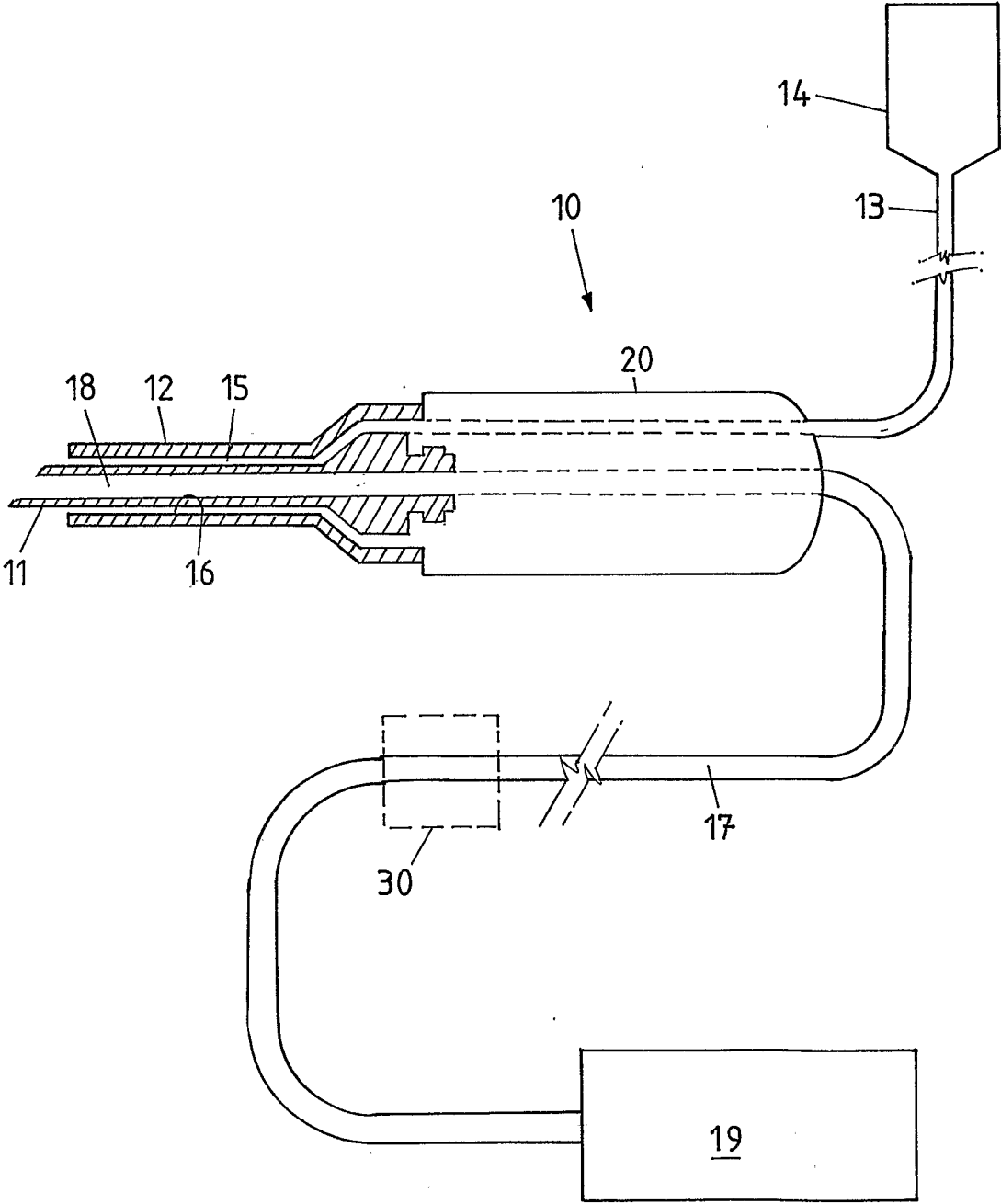


FIG.1

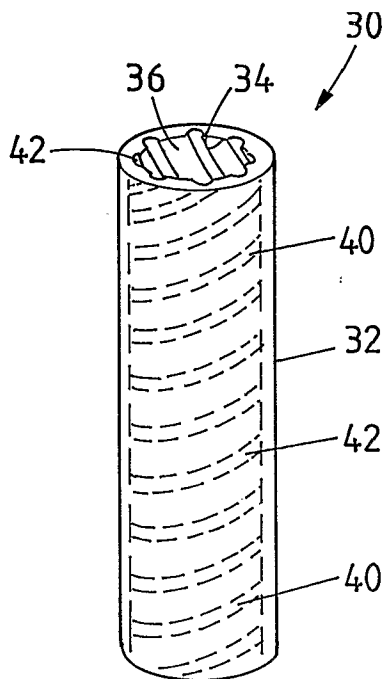


FIG. 2

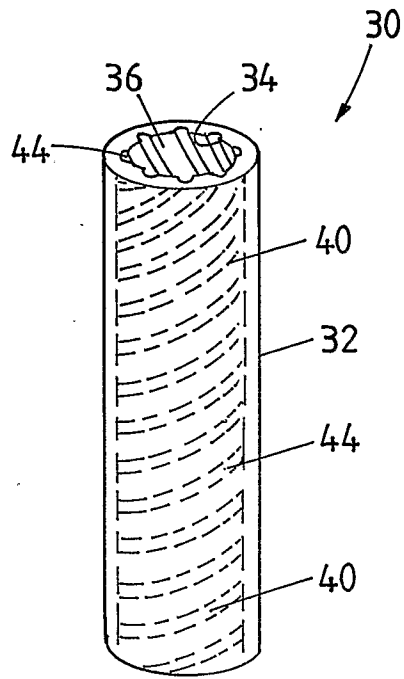


FIG. 3

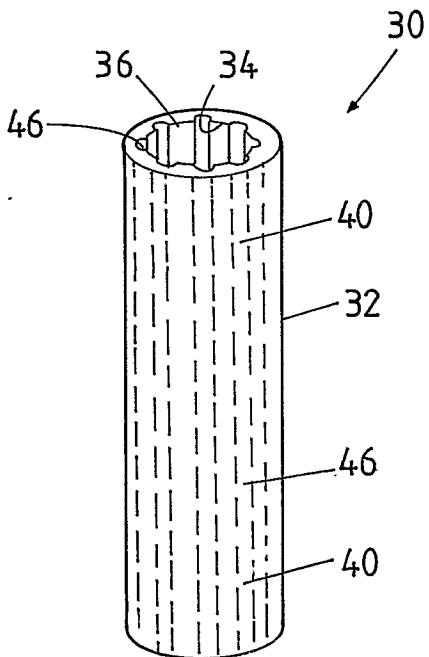


FIG. 4

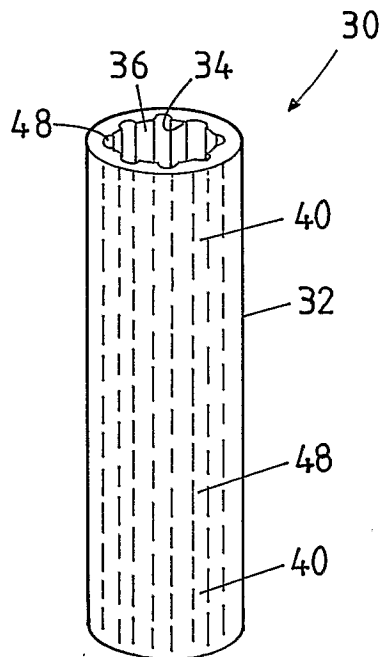


FIG. 5

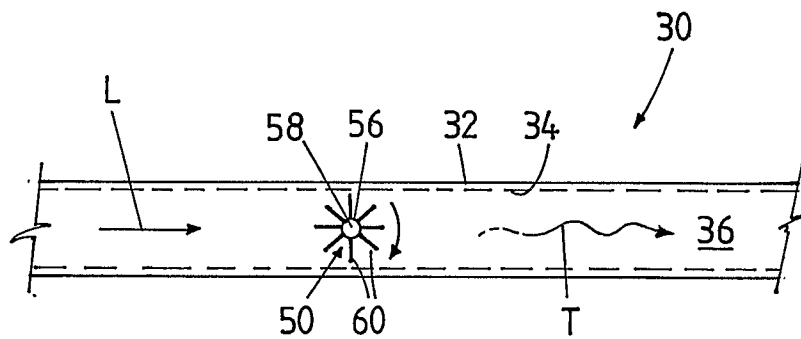


FIG. 6

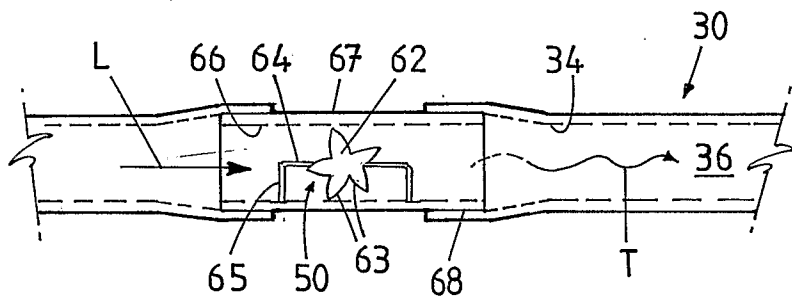


FIG. 7

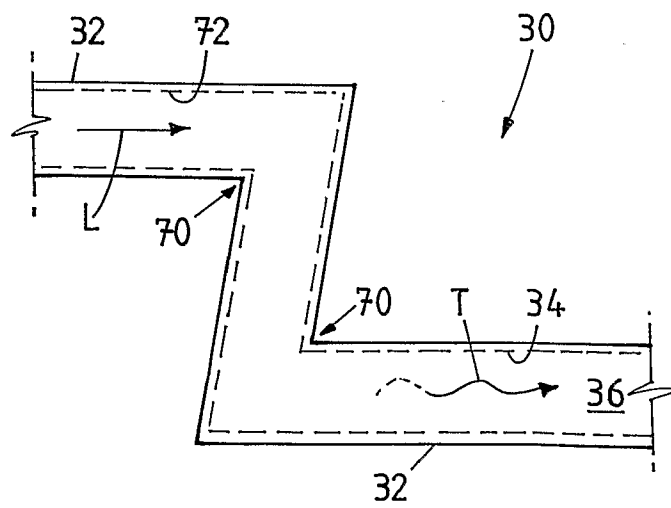


FIG. 8

FIG.9

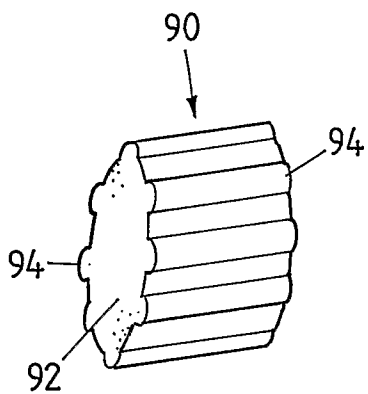
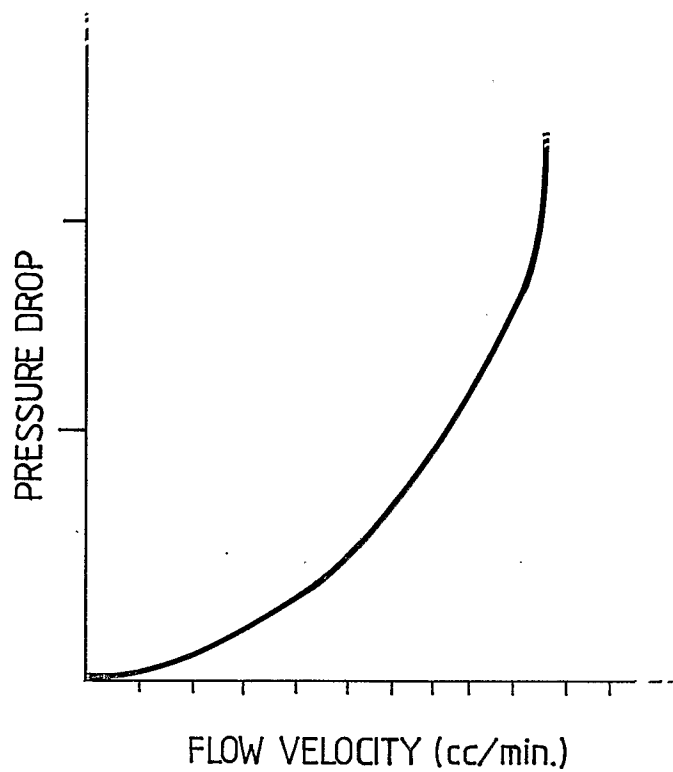


FIG. 10a.

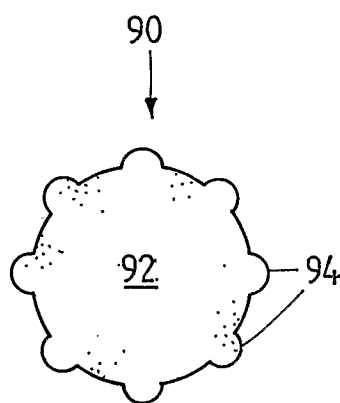


FIG. 10b.

INTERNATIONAL SEARCH REPORT

International application No.

PCT/AU03/00703

A. CLASSIFICATION OF SUBJECT MATTERInt. Cl. ⁷: A61M 1/00

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

DWPI IPC A61M, A61B, A61C, A61F and keywords (aspir, tub, flow, disrupt) and like terms

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 2002/0022810 A1 (URICH) 21 February 2002 paragraphs 21, 22 Fig 2	11, 12, 19, 22
X, P	US 2002/0128560 A1 (URICH) 12 September 2002 whole document	12, 19
A	WO 01/76681 A1 (CIRCUIT TREE MEDICAL INC) 18 October 2001 whole document	

☒ Further documents are listed in the continuation of Box C☒ See patent family annex

* Special categories of cited documents:

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier application or patent but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art

"&" document member of the same patent family

Date of the actual completion of the international search

5 August 2003

Date of mailing of the international search report

11 AUG 2003

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INTERNATIONAL SEARCH REPORT

International application No.

PCT/AU03/00703

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US 4041947 A (WEISS et al) 16 August 1977 whole document	
A	US 5167620 A (URECHE et al) 1 December 1992 column 5, lines 27-35, Fig. 1	
A	EP 0873762 A2 (JOHNSON & JOHNSON PROFESSIONAL INC) 28 October 1998 abstract	
A	US 5899884 A (COVER et al) 4 May 1999 abstract	

Box I Observations where certain claims were found unsearchable (Continuation of item 2 of first sheet)

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☐ Claims Nos :
because they relate to subject matter not required to be searched by this Authority, namely:
2. ☐ Claims Nos :
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:
3. ☐ Claims Nos :
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a)

Box II Observations where unity of invention is lacking (Continuation of item 3 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

See extra sheet.

1. ☐ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims
2. ☒ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest ☐ The additional search fees were accompanied by the applicant's protest.
☐ No protest accompanied the payment of additional search fees.

Supplemental Box

(To be used when the space in any of Boxes I to VIII is not sufficient)

Continuation of Box No: II

The international application does not comply with the requirements of unity of invention because it does not relate to one invention or to a group of inventions so linked as to form a single general inventive concept. In coming to this conclusion the International Searching Authority has found that there are two different inventions as follows:

1. Claims 1 to 7, 11 to 19, and 22 are directed to aspiration tubing comprising features involving the shape or configuration of the tubing to enhance resistance to fluid flow. It is considered that the shape or configuration of the tubing comprises a first "special technical feature".
2. Claims 8 to 10, 20 and 21 are directed to aspiration tubing comprising a freely moving object disposed within a lumen to disrupt the formation of laminar flow and enhance resistance to fluid flow through the lumen. It is considered that the freely moving object disposed within a lumen comprises a second "special technical feature".

These groups are not so linked as to form a single general inventive concept, that is, they do not have any common inventive features, which define a contribution over the prior art. The common concept linking together these groups of claims is that they have a means within the lumen to enhance resistance to fluid flow through the lumen. However this concept is not novel in the light of US 2002/0022810 A1 URICH. Therefore these claims lack unity a posteriori.

However since all these inventions share the same classification under the IPC they could be searched together without effort which would warrant an additional fee. Therefore all the inventions have been searched without extra charge.

INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No.

PCT/AU03/00703

This Annex lists the known "A" publication level patent family members relating to the patent documents cited in the above-mentioned international search report. The Australian Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

Patent Document Cited in Search Report				Patent Family Member			
US	2002022810	BR	200006376	CA	2426615	WO	2003030717
		AU	200151298	CA	2372277	EP	1189657
		US	6478781	WO	200176681		
US	2002128560	CA	2394177	EP	1281377	JP	2003102767
WO	200176681	AU	200151298	BR	200006376	CA	2372277
		EP	1189657	US	6478781	US	2002022810
		CA	2426615	WO	2003030717		
US	4041947	CA	1068572	CA	1068574	US	3902495
US	5167620	EP	625916	US	5106367	AU	21984/92
		WO	9315776				
EP	873762	CA	2235200	JP	11033108	US	6126628
US	5899884	NONE					
END OF ANNEX							